Pilot Study of DEXA Estimation in Pediatric Sick Patients (January-May 2017) At SGT Medical College Gurgaon (Haryana): How To Achieve The Best Results For The Management?

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ABSTRACT

Background: Osteoporosis can be seen in children because of chronic medical illness or medication. The evaluation leads to decrease risk of fractures by early intervention and management. A radiologist has to play an important role in the interpretation of the DEXA (Dual Energy X-ray Absorptiometery) examination results in pediatric age group. The role extends to the level of knowledge being statistician, orthopedic specialist and clinical pathologist. The various factors have been explored which affects the bone mineral density and have been compared with the available standard database. Bone mineral density (BMD) adds to detailed status of overall bone health. There have been many factors which affect the growing skeleton in children. The radiologist should know the technical aspect of acquiring the information. The interpretation is always based on the additional information other than DEXA scan. Osteoprosis is diagnosed with Z scoring and other biochemical evaluation parameters. T-scores in adults are totally different from the children group. Methods: Twenty pediatric patients (n=20) were subjected to DEXA examination from January to May 2017.All these children were from 10-19 years age group with mixed gender. These patients were of symptoms of common ailments as that of respiratory, renal, gastrointestinal and others. Results: Ten out of twenty (50%) cases were found to be having osteopenia or osteoporosis. The saensitivity of picking up the bone density was high in estimation of lumbosacral spine as compared to the femoral neck region. One case of mal absorption syndrome was exclusively having osteopenia (10%) and others were found to be having osteoporosis (90%). The common positive ailments were found to be that of tuberculosis (40%, malabsorption(30%), kidney disease(10%), diabetes mellitus (10%) and worms infestation(10%). Males were dominating in this group (72.7%) as compared to females (27.2%). Three (15%) were declared as of normal range. Conclusion: DEXA scan Z-scoring estimation is very sensitive method for diagnosing osteoporosis and osteopenia among pediatric age group. This is non invasive with minimal non-harmful radiation exposure. This is useful in the chronic ailments like tuberculosis, diabetes mellitus, malabsorption syndrome and kidney diseases. This is helpful in preventing the risks of fractures among these vulnerable patients.

Keywords: Osteoporosis; DEXA; BMD; Z score; T score.

INTRODUCTION

The incidence of osteoporosis can occur in children because of some primary bone disorders or secondary illness. The latter group includes those having indirect effects like immobility, anaemia, mal absorption and inflammation. Medication can also cause depletion of bone contents like that of steroids and chemotherapy. DEXA was introduced in 1987 and now it is widely used for the diagnosis. The interpretation of DEXA in children is quite challenging as there is continuous changing bone structure because of the ongoing development. The interpretation is in the form of numbers and the radiologist should have sufficient knowledge for the

clinical correlation. There are various factors which can influence the results like height, weight, body composition, gender, ethinicity and the physiological maturity. The patient nutritional and physical activity history is important for the evaluation

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Equipment and Procedure

All the cases were carried out on Hologen Discovery Wi (S/N 8835) model (Figure 1). The proper consent of the patient was taken .Lumbar region and femoral

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neck regions were included in the scanning.L1 to L4 vertebrae were taken into consideration. All the metal articles like buttons and clips etc. were removed before the test. The radiation dose of this test is 0.001 mSv which is equivalent to three hours of natural background radiation.



Figure 1: Hologen Discovery DEXA- scan machine with padded table top and overhead detector.

MATERIALS AND METHODS

Twenty pediatric patients (n=20) were subjected to DEXA examination at SGT Medical College Gurgaon from January to May2017 (Figure 2)7. These patients were of symptoms of common ailments as that of respiratory, renal, gastrointestinal and others (Figure 3). The group constituted of 11 male (55%) and 9 females(45%). The age ranged from 10-19 years (average 15.05 year). The weight ranged from 22-70 kg (average 43.4 kg). The height ranged from 131-170 cm. (average 149.15 cm) (Table1). Two sites were chosen for the examination as that of lumbosacral spine and either of femoral necks. The proper protocol was followed as described.

RESULTS

The estimation was evaluated as per WHO Zscoring, The values less than -2.5 was taken as that of osteopenia and more than that was counted among osteoporosis. Ten out of twenty (50%) cases were found to be having osteopenia or osteoporosis. The incidence of decreased BMD was more seen in male patients (40%) as compared to female (15%) group (Figure 5). The sensitivity of picking up the bone density was high in estimation of lumbosacral spine as compared to the femoral neck region. One case of mal absorption syndrome was exclusively having osteopenia (10%) and others were found to be having osteoporosis (90%). The common positive were found to be that of tuberculosis ailments (40%, malabsorption(30%), kidney

disease(10%),diabetes mellitus (10%) and worms infestations(10%) (Figure 4).Six were having

osteopenia (30%) and the ratio was equally distributed among both the sexes. Eleven (55%) the patients were having osteoporosis .Males were dominating in this group (72.7%) as compared to females (27.2%).Three (15%) were declared as of normal range.Tgiven in details in Table 2.he detailed stastics of the patients had been.

	Total Patients	Percentage
Male	11	55
Female	9	45

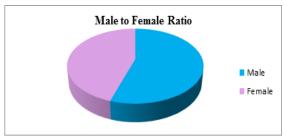


Figure 2: Male to female ratio of DEXA patients

	Normal	With ailments
Male Patients	3	8
Female Patients	7	2

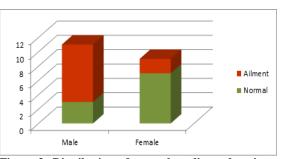


Figure 3: Distribution of normal to diseased patients with sex linkage.

Patient No.	Weight (kg)	Height	Age
		(cm)	(Years)
1	25	131	10
2	40	166	12
3	50	170	16
4	48	148	17
5	22	104	13
6	33	142	11
7	48	153	14
8	47	149	16
9	44	151	13
10	42	159	17
11	45	160	14
12	70	162	17
13	64	159	19
14	36	150	17
15	25	134	12
16	31	131	16
17	40	145	13
18	46	149	18
19	63	168	19
20	49	152	17

Average Weight = 43.4 kg Average height = 149.15 cm

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Average Age =15.05 years

Table 1: Detailed statistics of the patients included in the study

the stu	uy	
S.No.	Ailment	Number of patients
1	Tuberculosis	4
2	Worm Infestation	1
3	Malabsorption	3
4	Kidney disease	1
5	Diabetes Mellitus	1

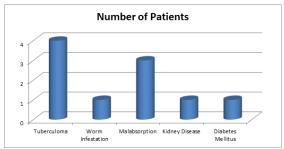


Figure 4: Detailed of the ailments found in the group which was included in the study.

	Male	Female
Osteopenia	3	3
Osteoporosis	8	3
Normal	0	3

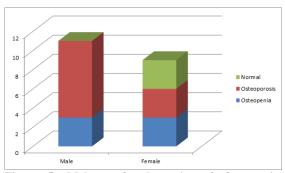


Figure 5: Male to female ratios of Osteopenia, Osteoporosis and Normal patients.

Table	Table 2: Chart showing the details of all the pediatric DEXA cases during the period of Jan 2017-May 2017										17-May 2017.	
S.No	Nam e	Age	Weigh t (Kg)	Heigh t (Cm)	Complain	Regio n	Area (Cm2	Bm c (G)	Bmd (G/Cm 2)	T- Scor e	Z- Scor e	Remarks
1.	S	10/ M	25	131	Multiple Cerebral Tuberculomas On ATT X 6	Spine L1-L4 Femor	35.28 21.04	14.1 7 11.2	0.402	-6.3	-6.3	Tuberculosis. On calcium supplement
_					months	al Neck		9				
2.	L	12/ M	40	166	Mal absorption Syndrome	Spine L1-L4	52.49	37.9 2	0.722	-3.4	-3.4	On treatment with dietary
						Femor al Neck	30.60	26.5	0.867	-1.1	-1.1	regulation.
3.	S	16/ M	50	170	Cervical Lymphadenopath	Spine L1-L4	49.82	43.8 2	0.880	-1.9	-1.9	Under Follow up
					y discharging sinus on lrft posterior auricular region	Femor al Neck	32.66	33.5 5	1.027	0.0	0.0	
4.	N	17/F	48	148	Renal Calculus, No systemic	Spine L1-L4	45.86	48.5 6	1.059	0.1	0.3	Normal Follow up
					Disease	Femor al Neck	24.27	20.8	0.858	-0.7	-0.7	
5.	D	13/F	22	104	Recurrent loose motions with mal	Spine L1-L4	35.08	19.7 1	0.562	-4.8	-4.8	Dietary control and
					absorption syndrome.On treatment	Femor al Neck	26.99	20.0	0.743	-1.9	-1.9	calcium supplement
6.	D	11/ M	33	142	Multiple mesenteric node	Spine L1-L4	35.30	19.6 7	0.557	-4.9	-4.9	.Tuberculosis On calcium
					with pain abdomen- 4-5 yrs	Femor al Neck	22.50	15.2	0.676	-2.4	-2.4	supplement
7.	S	14/F	48	153	Pain Epigastric-2 yrs	Spine L1-L4	49.67	44.4 0	0.894	-1.4	-1.2	Normal Follow up
						Femor al Neck	28.39	26.8 2	0.945	0.0	0.0	
8.	M	16/F	47	149	Pain Abdomen (? Left Ureteric	Spine L1-L4	49.15	40.4 8	0.824	-2.0	-1.9	Normal Follow up
					calculus)	Femor al Neck	29.32	22.2 4	0.759	-1.5	-1.5	
9.	U	13/F	44	151	Pain Abdomen- 2	Spine	49.66	38.7	0.781	-2.4	-2.3	Normal

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Femor al Neck Neck		I	1			yrs	L1-L4	1	6	1			Follow up
10. S						J15		31.97		0.783	-1.3	-1.3	_ ronow up
10. S									3				
Next	10	-	17/	10	150	NY 1 2		50.00	24.2	0.672	2.0	2.0	
Femor Sect Femo	10.	S		42	159	1		50.90		0.672	-3.8	-3.8	On treatment
1.1			141			syndrome		38.03		0.825	-1.4	-1.4	
11. R							al		7				
M					1 -0								
Second S	11.	R		45	160			55.78		0.727	-2.9	-2.7	
12. R C 17/ 70 162 Mild Spleenomegaly Femor 45.14 44.9 0.996 0.4 0.4 0.4 1.14 1.0 0.996 0.4 0.4 0.4 1.14 1.0 0.996 0.4 0.4 0.4 1.14 1.0 0.996 0.4 0.4 0.4 1.14 1.0 0.996 0.4 0.4 0.4 1.14 1.0 0.996 0.4 0.4 0.4 1.14 0.996 0.4 0.4 0.4 1.14 0.996 0.4 0.4 0.4 1.14 0.996 0.4 0.4 0.4 1.14 0.996 0.4 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 0.4 1.14 0.996 0.4 0.4 1.14 0.996 0.4 0.4 0.4 1.14 0.996 0.4 0.4 0.4 1.14 0.996 0.4 0.4 0.4 1.14 0.996 0.4 0.4 0.4 1.14 0.996 0.4 0.4 0.4 1.14 0.996 0.4 0.4 0.4 1.14 0.996 0.4 0.4 0.4 1.14 0.996 0.4 0.			IVI					36.20		0.907	-0.3	-0.3	
12.						• 1		20.20		0.507	0.5	0.0	
12. R C 17/ N 13/ M 162 Mild Spleenomegaly Spine S5.15 47.2 0.856 -1.7 -1.6 Normal Follow up						1.1	Neck						
13. R 19/F 64 159 Renal Stones Spine 11-14 4 44.9 0.996 0.4 0.4 0.4 1.4 1.4 1.4 1.4 1.4 1.5	12	P.C	17/	70	162		Cnino	55 15	47.2	0.956	1.7	1.6	Normal
13. R 19/F 64 159 Renal Stones Spine 56.25 50.5 0.899 -1.3 -1.2 Normal Follow up	12.	RC		70	102			33.13		0.830	-1./	-1.0	
13. R 19/F 64 159 Renal Stones Spine 1.1-1.4 Femor 29.01 28.2 0.975 0.3 0.3 0.3 0.3 0.3 0.4 0.5 0.899 1.3 0.3			1.1			Spicenomegary		45.14		0.996	0.4	0.4	_ rono w up
13. R 19/F 64 159 Renal Stones Spine L1-1.4 C1-1.4 Follow up Follow									4				
14. A 17/ 36 150 Recurrent loose motions and pain abdomen Spine 29.01 28.2 0.975 0.3 0.3 0.3 0.3 14. A 17/ 36 150 Recurrent loose motions and pain abdomen Spine 28.60 18.5 0.649 -2.0 -1.5 0.0 Follow up 15. J 12/ 25 134 Multiple mesenteric lymph modes are present, pain abdomen(epigastr ic) Spine 36.78 23.3 0.635 -4.1 -4.1 Tuberculosis on ATT and calcium supplement Spine 21.67 15.4 0.712 -2.1 -2	1.2		10/5	6.1	150	D 10:		56.25	50.5	0.000	1.0	1.2	N. 1
14. A 17/ 36	13.	R	19/F	64	159	Renal Stones		56.25		0.899	-1.3	-1.2	
14. A 17/ 36								29.01		0.975	0.3	0.3	1 onow up
14. A									9				
M					1.50								
Spine Spin	14.	A		36	150			42.67		0.674	-2.8	-1.8	On Follow up
15.			IVI					28.60		0.649	-2.0	-1.5	
15.								20.00		0.0.5	2.0	1.0	
M													
16. A 16/F 31 131 Pain abdomen Diagnosed as that of worms infestation Spine L1-L4 Femor al Neck Spine al	15.	J		25	134			36.78		0.635	-4.1	-4.1	
Pain abdomen(epigastric) Pain abdomen Pain ab			IVI					21.67		0.712	-2.1	-2.1	
16. A 16/F 31 131 Pain abdomen Diagnosed as that of worms infestation Pain abd men Diagnosed as that of worms infestation Pain abd men Diagnosed as that of worms infestation Pain abd with Pain abd. Wit								21.07		0.712	2.1	2.1	
16. A 16/F 31 131							Neck						
17. N	16	_	16/E	21	121	, ,	Cnino	27.60	10 1	0.656	2.6	2.2	On diotory and
17. N 13/ M 40 145 Pain abd. With yellowish urine USG- Multiple subcentrimetric mesenteric lymph nodes 18. A 18/F 46 149 Rt. Lumbar region pain- 3-4 days 27.5 19. 19/ M 1	10.	Α	10/1	31	131			27.09		0.030	-3.0	-3.3	
17. N 13/ M 145 Pain abd. With yellowish urine USG- Multiple subcentrimetric mesenteric lymph nodes 18. A 18/F 46 149 Rt. Lumbar region pain- 3-4 days Remore 27.94 25.4 0.912 -0.2 -0.2 -0.2								19.11		0.819	-1.0	-1.0	
17. N 13/ M 40 145 Pain abd. With yellowish urine USG-Multiple subcentrimetric mesenteric lymph nodes 18. A 18/F 46 149 Rt. Lumbar region pain- 3-4 days Remore all Neck 149 Rt. Lumbar region pain- 3-4 days 152 Loss of appetite- 5-6 months 17/F 49 152 Loss of appetite- 5-6 months 18/F 17/F 18/F 18/F						infestation			4				
17. N							Neck						
M	17.	N	13/	40	145	Pain abd. With	Spine	42.93	28.1	0.656	-3.6	-3.4	
Subcentrimetric mesenteric lymph nodes Spine Section Spine Section Spine Section Spine Section Spine Section Spine Section Secti			M										
18. A 18/F 46 149 Rt. Lumbar region pain- 3-4 days Spine L1-L4 Femor al Neck Spine L1-L4 Spine Spine								29.88		0.765	-1.5	-1.5	supplement
18. A 18/F 46 149 Rt. Lumbar region pain- 3-4 days Spine L1-L4 Femor al Neck Spine L1-L4 Spine L1-L5 Spine L1-L4 Spine									6				
Tegion pain- 3-4 days California Calif							IVCCK						
19. J	18.	A	18/F	46	149	Rt. Lumbar		54.42		0.922	-1.1	-1.0	
19.								27.04		0.012	0.2	0.2	Follow up
19. J 19/ 63 168 Dyspepsia with Malabsorption syndrome, USG-Cholelithiasis with chronic cholecystitis 20. N 17/F 49 152 Loss of appetite-5-6 months 154.92 155 1.066 1.067 1						uays		27.94		0.912	-0.2	-0.2	
19.									U	1			
Syndrome, USG- Cholelithiasis with chronic cholecystitis	19.	J		63	168		Spine	51.09		0.863	-2.1	-2.1	
20. N 17/F 49 152 Loss of appetite-5-6 months Spine L1-L4 Spine L1-L4 Seminary Spine L1-L4 Seminary Spine L1-L4 Seminary Spine L1-L4 Seminary Spine Sp			M			1		22.05		0.010	1	1.7	up
Cholelithiasis with chronic cholecystitis							-	33.86		0.813	-1.5	-1.5	
20. N 17/F 49 152 Loss of appetite-5-6 months Spine L1-L4									3	1			
20. N 17/F 49 152 Loss of appetite-5-6 months Spine L1-L4 Spine L1-L4 Femor 36.08 31.6 0.878 -0.5 -0.5 Follow up													
5-6 months L1-L4 3 Follow up Femor 36.08 31.6 0.878 -0.5 -0.5	20	N.	10.7	40	1.52		G :	5402	50.5	1000	0.2	0.2	NT 1
Femor 36.08 31.6 0.878 -0.5 -0.5	20.	IN	1 //F	49	152			54.92		1.066	0.2	0.3	
al 9						o monuis		36.08		0.879	-0.5	-0.5	- 1 ono up
								30.00		0.076	-0.5	-0.5	

DISCUSSION

The evaluation by DEXA becomes more significant in following situations:

When more than two long bone fractures present before 10 years of age.

When more than three fractures are present upto 19 years of age.

Lastly when Z score is >-2.0

Pediatric DEXA scan interpretation requires a great expertise because of the growing skelton. There are many other factors, like type of indication, volume

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versus area, technical factors and protocol in newborns, to be kept in mind for the accurate results.. Bone Mineral Density for the region of interest (ROI) is calculated by dividing the Bone Mineral Contents (BMC) in grams by Bone area (BA) in squared centimeters. This is always compared with the reference standard.BA of ROI is always taken in two dimensions comparing to the actual data which denotes three dimensional region [1]. The volume of the bone keeps on changing in the growing skeleton so the frequent follow ups are suggested [2].Lumbar region and hip joints are selected for the scan.L1 to L4 vertebrae are properly centered with inclusion of minimal soft tissue. Hip joint should not have overlapping trochanters,neck and acetabulum regions.





Figure 6: DEXA scan protocol. Lumbar spine (L1-L4) and hip joint should be centered as shown in the diagram.

This also should not include any tubes, jewellary or prosthetics.Adult algorithm always overestimates as BMD in pediatric patient is very low. Sometimes thoracic spine is included in the study and this is the best way to pick up collapsed vertebra [3,4]. Ashes bone is the real assessment of the BMC but it represents 7-9% of the bone ash value. There are variations due to wrong positioning and motion artifacts. Calibration is required with the phantom for the daily quality control scan. Least significant change (LSC) should be observed in follow up as per the age in 6-12 months.BMC and BA are closely related weight and height in the new born [5,6].

There is strong indication for DEXA in children who are on corticoids for more than two months. The indication for other conditions like chronic inflammatory disease, hypogonadism, prolonged immobilization, low trauma fractures and where radiographs gives clue for osteoporosis. Few of the syndromes and genetic conditions also requires this evaluation like Ehlers-Danlos , Fibrous dysplasia, Homocystinuria, Hypophosphatasia, idiopat hic hypercalciuria, Marfan's syndrome, osteogenesis imperfect and Menke's kinky hair syndrome.

In newborns it is area (aBMD) rather than volumetric data (vBMD appropriate. Osteopenia and osteoporosis are defined as per WHO criteria if T-score < -1 and < -2.5 respectively. Z score in children show low mean BMD values as bone peak has yet to be achieved. In the report "low bone density" can be reported to avoid any confusion. This increase rapidly in puberty. Other dependent factors are ethinicity, gender, age and physiological maturity levels. These have to be compared with pediatric normative database.Gafni and Baron (2004) reported half of the cases of misdiagnosed osteoporosis as given by by T-score [7,8]. Tanner stage or gynecologic age have been included for the correct results. Crabtree et al (2004) proposed lean total body mass (LTM) algorithm in relation to height and BMC [9]. American college of Radiology and Americal college of Rheumaatology recommend DEXA for the children but Society of Pediatric Radiology and American academy of Pediatrics have not cleared it so far. The radiation dose to the patient is negligible and even less than the normal background radiation dose (7micro Sv). This has got no biological effect on the organs .Njhe et al (1997) had demonstrated that scatter dose is less than 1 micro mSv even the operator sitting at one meter distance without shielding. Thus annual dose for the operator is approximate 0.4 mSv with workload of sixteen patients per day [10].

CONCLUSION

DEXA scan Z-scoring estimation is very sensitive method for diagnosing osteoporosis and osteopenia among pediatric age group. This is non invasive with minimal non-harmful radiation exposure. The incidence is slightly more among male sick patients as compared to the female group. The incidence has also been noticed higher among the patients who are on long term medication. This is useful in the chronic ailments like tuberculosis, diabetes mellitus, malabsorption syndrome and kidney diseases. This is helpful in preventing the risks of fractures among these vulnerable patients.

Acknowledgement

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Consent

Written consent of the parents was taken before performing the DEXA scan test.

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